EGG ALLERGEN MIX- adenosinum triphosphoricum dinatrum, l-asparagine (monohydrate), l-phenylalanine, quercetin, ileum (suis), jejunum (suis), stomach (suis), ascorbicum acidum, calcarea carbonica, ferrum iodatum, tetracycline, egg (hen whole), proteus (morgani) liquid
Deseret Biologicals, Inc.

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

DRUG FACTS:

ACTIVE INGREDIENTS:

Adenosinum Triphosphoricum Dinatrum 4X, L-Asparagine (Monohydrate) 6X, 10X, 30X, 200X, L-Phenylalanine 6X, 10X, 30X, 200X, Quercetin 6X, 10X, 30X, 200X, Ileum (Suis) 8X, Jejunum (Suis) 8X, Stomach (Suis) 8X, Ascorbicum Acidum 6C, 30C, Calcarea Carbonica 6C, 30C, Ferrum Iodatum 6C, 30C, Tetracycline 6C, 30C, Egg (Hen Whole) 28C, Proteus (Morgani) 30C, 200C.

INDICATIONS:

For the temporary relief of symptoms related to allergies to eggs in foods and supplements, including skin inflammation, hives, runny nose, sneezing, stomach cramps, nausea, vomiting, coughing.**

**These statements are based upon homeopathic principles. They have not been reviewed by the Food and Drug Administration.

WARNINGS:

Keep out of reach of children. In case of overdose, contact a physician or Poison Control Center right away.

If pregnant or breast-feeding, ask a health professional before use.

Tamper seal: "Sealed for Your Protection." Do not use if seal is broken or missing.

KEEP OUT OF REACH OF CHILDREN:

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

DIRECTIONS:

3-10 drops under the tongue, 3 times a day or as directed by a health professional. Consult a physician for use in children under 12 years of age.

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INACTIVE INGREDIENTS:

Demineralized Water, 25% Ethanol

QUESTIONS:

Dist. By: Deseret Biologicals, Inc.

469 W. Parkland Drive

Sandy, UT 84070 www.desbio.com

PACKAGE LABEL DISPLAY:

DES**BIO**

NDC 43742-1644-1

HOMEOPATHIC

EGG ALLERGEN

MIX

1 FL OZ (30 ml)

EGG ALLERGEN MIX

adenosinum triphosphoricum dinatrum, l-asparagine (monohydrate), l-phenylalanine, quercetin, ileum (suis), jejunum (suis), stomach (suis), ascorbicum acidum, calcarea carbonica, ferrum iodatum, tetracycline, egg (hen whole), proteus (morgani) liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:43742-1644
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ADENOSINE TRIPHOSPHATE DISODIUM (UNII: 5L51B4DR1G) (ADENOSINE TRIPHOSPHATE - UNII:8L70Q75FXE)	ADENOSINE TRIPHOSPHATE DISODIUM	4 [hp_X] in 1 mL
ASPARAGINE MONOHYDRATE (UNII: 2PD79VF521) (ASPARAGINE - UNII:5Z33R5TKO7)	ASPARAGINE MONOHYDRATE	6 [hp_X] in 1 mL
PHENYLALANINE (UNII: 47E5O17Y3R) (PHENYLALANINE - UNII:47E5O17Y3R)	PHENYLALANINE	6 [hp_X] in 1 mL
QUERCETIN (UNII: 9IKM0I5T1E) (QUERCETIN - UNII:9IKM0I5T1E)	QUERCETIN	6 [hp_X] in 1 mL
SUS SCROFA ILEUM (UNII: C998R1XSRA) (SUS SCROFA ILEUM - UNII:C998R1XSRA)	SUS SCROFA ILEUM	8 [hp_X] in 1 mL
SUS SCROFA JEJUNUM (UNII: TA501QD69R) (SUS SCROFA JEJUNUM - UNII:TA501QD69R)	SUS SCROFA JEJUNUM	8 [hp_X] in 1 mL
SUS SCROFA STOMACH (UNII: T0920P9Z9A) (SUS SCROFA STOMACH - UNII:T0920P9Z9A)	SUS SCROFA STOMACH	8 [hp_X] in 1 mL
ASCORBIC ACID (UNII: PQ6CK8PD0R) (ASCORBIC ACID - UNII:PQ6CK8PD0R)	ASCORBIC ACID	6 [hp_C] in 1 mL
OYSTER SHELL CALCIUM CARBONATE, CRUDE (UNII: 2E32821G6I) (OYSTER SHELL CALCIUM CARBONATE, CRUDE - UNII:2E32821G6I)	OYSTER SHELL CALCIUM CARBONATE, CRUDE	6 [hp_C] in 1 mL
FERROUS IODIDE (UNII: F5452U54PN) (FERROUS IODIDE - UNII:F5452U54PN)	FERROUS IODIDE	6 [hp_C] in 1 mL
TETRACYCLINE (UNII: F8VB5M810T) (TETRACYCLINE - UNII:F8VB5M810T)	TETRACYCLINE	6 [hp_C] in 1 mL
EGG (UNII: 291P45F896) (EGG - UNII:291P45F896)	EGG	28 [hp_C] in 1 mL
PROTEUS MORGANII (UNII: 56X6LID5ZY) (PROTEUS MORGANII - UNII:56X6LID5ZY)	PROTEUS MORGANII	30 [hp_C] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43742-1644-1	30 mL in 1 PACKAGE; Type 0: Not a Combination Product	05/11/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		05/11/2020	

Labeler - Deseret Biologicals, Inc. (940741853)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-1644) , api manufacture(43742-1644) , label(43742-1644) , pack(43742-1644)

Revised: 1/2024 Deseret Biologicals, Inc.